AMENDMENTS TO THE CLAIMS

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1. (Currently Amended) A triazolopyrimidine compound of the formula I

in which the substituents are as defined below:

 R^1 is C_2 - C_{12} -alkenyl or C_2 - C_{12} -alkynyl, where the carbon chains are unsubstituted or carry one to three identical or different groups R^a and/or R^b :

or

 C_1 - C_{14} -alkyl, C_1 - C_{12} -alkoxy- C_1 - C_{12} -alkyl, C_1 - C_6 -alkoxy- C_2 - C_{12} -alkynyl, where the carbon chains carry one to three identical or different groups R^a ;

 R^a is halogen, cyano, nitro, hydroxyl, C_1 - C_6 -alkylthio, C_3 - C_{12} -alkenyloxy, C_3 - C_{12} -alkynyloxy, or

C₃-C₆-cycloalkyl which may carry one to four identical or different groups R^b;

R^b is C₁-C₄-alkyl, cyano, nitro, hydroxyl, C₁-C₆-alkoxy, C₁-C₆-alkylthio, C₃-C₆-alkenyloxy and C₃-C₆-alkynyloxy;

where the carbon chains of the groups Ra for their part may be halogenated;

- R^2 is C_1 - C_{12} -alkyl, C_2 - C_{12} -alkenyl or C_2 - C_{12} -alkynyl, where the carbon chains are substituted by one to three groups R^c :
 - R^c is cyano, nitro, hydroxyl; or C₃-C₆-cycloalkyl which may carry one to four identical or different groups C₁-C₄-alkyl, halogen, cyano, nitro, hydroxyl, C₁-C₆-alkoxy, C₁-C₆-alkylthio, C₃-C₆-alkenyloxy or C₃-C₆-alkynyloxy.
- 2. (Original) The compound of the formula I according to claim 1 in which
 - R^1 is C_1 - C_{14} -haloalkyl, C_1 - C_{12} -haloalkoxy- C_1 - C_{12} -alkyl, C_1 - C_{12} -alkoxy- C_1 - C_{12} -haloalkyl, C_2 - C_{12} -alkenyl, C_2 - C_{12} -haloalkenyl, C_2 - C_{12} -alkynyl or C_2 - C_{12} -haloalkynyl, where the carbon chains may carry one to three groups R^a :
 - R^a is cyano, nitro, hydroxyl, C₁-C₆-alkoxy, C₁-C₆-alkylthio, C₃-C₁₂-alkenyloxy, C₃-C₁₂-alkynyloxy, or

C₃-C₆-cycloalkyl which may carry one to four identical or different groups;

R^b is C₁-C₄-alkyl, cyano, nitro, hydroxyl, C₁-C₆-alkoxy, C₁-C₆-alkylthio, C₃-C₆-alkenyloxy and C₃-C₆-alkynyloxy

where the carbon chains of the groups R^a for their part may be halogenated.

- 3. (Original) The compound of the formula 1 according to claim 1 or 2 in which
 - R^2 is C_1 - C_{12} -alkyl, C_2 - C_{12} -alkenyl or C_2 - C_{12} -alkynyl, where the carbon chains may be substituted by one to three groups R^c :
 - R^c is cyano, nitro, hydroxyl; or C₃-C₆-cycloalkyl which may carry one to four identical or different groups C₁-C₄-alkyl, halogen, cyano, nitro, hydroxyl, C₁-C₆-alkoxy, C₁-C₆-alkylthio, C₃-C₆-alkenyloxy or C₃-C₆-alkynyloxy.
- 4. (Previously Presented) The compound of the formula I according to claim 1 in which
 - R¹ is C₁-C₁₄-alkyl, where the carbon chains carry one to three identical or different groups cyano or halogen.
- 5. (Previously Presented) The compound of the formula I according to claim 1 in which

 R^1 is C_2 - C_{12} -alkenyl or C_2 - C_{12} -alkynyl, where the carbon chains are unsubstituted or

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6. (Previously Presented) The compound of the formula I according to claim 1 in which R¹ and R² together do not have more than 14 carbon atoms.

carry one to three identical or different groups R^a and/or R^b.

(Previously Presented) The compound of the formula I according to claim 1 in which R¹ is 7. chloromethyl, bromomethyl, dichloromethyl, trichloromethyl, fluoromethyl, difluoromethyl, trifluoromethyl, chlorofluoromethyl, dichlorofluoromethyl, chlorodifluoromethyl, 1-chloroethyl, 1-bromoethyl, 1-fluoroethyl, 2-fluoroethyl, 2,2difluoroethyl, 2,2,2-trifluoroethyl, 2-chloro-2-fluoroethyl, 2-chloro-2,2-difluoroethyl, 2,2dichloro-2-fluoroethyl, 2,2,2-trichloroethyl, pentafluoroethyl, 1,1,1-trifluoroprop-2-yl, 1chloropropyl, 1-fluoropropyl, 3-chloropropyl, 3-fluoropropyl, 3,3,3-trifluoropropyl, 1chlorobutyl, 1-fluorobutyl, 4-chlorobutyl, 4-fluorobutyl, 4,4,4-trifluorobutyl, 1chloropentyl, 1-fluoropentyl, 5,5,5-trifluoropentyl, 5-chloropentyl, 5-fluoropentyl, 1chlorohexyl, 1-fluorohexyl, 6-chlorohexyl, 6-fluorohexyl, 6,6,6-trifluorohexyl, 1chloroheptyl, 1-fluoroheptyl, 7-chloroheptyl, 7-fluoroheptyl, 7,7,7-trifluoroheptyl, 1chlorooctyl, 1-fluorooctyl, 8-fluorooctyl, 8,8,8-trifluorooctyl, 1-chlorononyl, 1fluorononyl, 9-fluorononyl, 9,9,9-trifluorononyl, 9-chlorononyl, 1-fluorodecyl, 1chlorodecyl, 10-fluorodecyl, 10,10,10-trifluorodecyl, 10-chlorodecyl, 1-chloroundecyl, 1fluoroundecyl, 11-chloroundecyl, 11-fluoroundecyl, 11,11,11-trifluoroundecyl, 1-chloro-

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dodecyl, 1-fluorododecyl, 12-fluorododecyl or 12,12,12-trifluorododecyl.

- 8. (Previously Presented) The compound of the formula I according to claim 1 in which R² is methyl, ethyl, isopropyl, n-propyl or n-butyl.
- 9. (Original) The compound of the formula I according to claim 1:

6-(3-bromopropyl)-5-ethyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

6-(3-chloropropyl)-5-ethyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

6-(7-amino-5-ethyl-[1,2,4]triazolo[1,5-a]pyrimidin-6-yl)-hexanenitrile;

6-(7-amino-5-propyl-[1,2,4]triazolo[1,5-a]pyrimidin-6-yl)-hexanenitrile;

5-ethyl-6-hex-5-enyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

6-hex-5-enyl-5-methyl-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine;

5-methyl-6-(5,6,6-trifluor ohex-5-enyl)-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine.

10. (Withdrawn) A process for preparing compounds of the formula I according to claim 1 wherein β-ketoesters of the formula II,

$$RO$$
 R^{1}
 R^{2}
 R^{2}
 R^{2}

in which R is C₁-C₄-alkyl are reacted with 3-amino-1,2,4-triazole of the formula III

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$$N \longrightarrow NH_2$$

to give 7-hydroxytriazolopyrimidines of the formula IV

$$\begin{array}{c|c}
 & \text{OH} \\
 & \text{N} \\
 & \text{N} \\
 & \text{N} \\
 & \text{R}^2
\end{array}$$

which are halogenated to give compounds of the formula V

$$\begin{array}{c|c}
 & \text{Hal} \\
 & \text{N} \\
 & \text{N} \\
 & \text{N} \\
 & \text{N} \\
 & \text{R}^2
\end{array}$$

in which Hal is chlorine or bromine and V is reacted with ammonia.

11. (Withdrawn) A process for preparing compounds of the formula I according to claim 1 wherein acylcyanides of the formula VI,

$$\begin{array}{cccc}
 & & & VI \\
 & & & & VI \\
 & & & & & VI
\end{array}$$

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are reacted with 3-amino-1,2,4-triazole of the formula III.

12. (Withdrawn) A compound of the formula IV or V according to claim 10.

13. (Withdrawn) A process for preparing compounds of the formula I according to claim 1 in which R¹ is halogen-substituted C₁-C₁₄-alkyl, C₁-C₁₂-alkoxy-C₁-C₁₂-alkyl, C₂-C₁₂-alkenyl or C₂-C₁₂-alkynyl, by halogenating triazolopyrimidines of the formula VII,

$$N - N$$
 R
 $N - N$
 R^2
 $N - N$
 R^2

in which R is C_1 - C_{14} -alkyl, C_1 - C_{12} -alkoxy- C_1 - C_{12} -alkyl, C_2 - C_{12} -alkenyl, C_2 - C_{12} -alkynyl, where the carbon chains may carry one to three groups R^a as set forth in claim 1, using a halogenating agent in the presence of a free-radical initiator or an acid.

- 14. (Previously Presented) A fungicidal composition comprising a solid or liquid carrier and a compound of the formula I according to claim 1.
- 15. (Withdrawn) Seed comprising a compound of the formula I according to claim 1 in an amount of 1 to 1000 g per 100 kg.
- 16. (Withdrawn) A method for controlling phytopathogenic harmful fungi wherein the fungi or the materials, plants, the soil or seed to be protected against fungal attack are treated with an effective amount of a compound of the formula I according to claim 1.

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